Regulation of Phosphorylation of the GluR1 AMPA Receptor in the Neostriatum by Dopamine and Psychostimulants *In Vivo*

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The activation of cAMP-dependent protein kinase regulates the physiological activity of AMPA-type glutamate receptors. In this study, phosphorylation of the AMPA receptor subunit GluR1 at Ser⁸⁴⁵ was increased in neostriatal slices by activation of D1-type dopamine receptors and by inhibitors of protein phosphatase 1/protein phosphatase 2A. In contrast, Ser⁸³¹, a residue which, when phosphorylated by protein kinase C or calcium/calmodulin-dependent kinase II, increases AMPA receptor channel conductance, was unaffected by either D1 or D2 receptor agonists in neostriatal slices. The phosphorylation of

Ser⁸⁴⁵, but not Ser⁸³¹, was strongly increased in neostriatum *in vivo* in response to the psychostimulants cocaine and methamphetamine. The effects of dopamine and psychostimulants on the phosphorylation of GluR1 were attenuated in dopamine and cAMP-regulated phosphoprotein *M_r* 32 kDa (DARPP-32) knock-out mice. These results identify DARPP-32 and AMPA-type glutamate receptors as likely essential cellular effectors for psychostimulant actions.

Key words: dopamine; DARPP-32; methamphetamine; cocaine; protein phosphatase 1; D1 receptor; protein kinase A

Ionotropic glutamate receptors function as ligand-gated ion channels and are believed to play a role in certain forms of learning and memory (Bliss and Collingridge, 1993; Linden, 1994), in the development of psychomotor stimulant sensitization (Wolf, 1998) and in various forms of neurodegenerative disease (Lipton and Rosenberg, 1994). They are classified by their physiological and pharmacological characteristics as members of AMPA, kainate, or NMDA receptor subclasses (Monaghan et al., 1989). AMPA-type glutamate receptors are abundantly expressed on dendritic processes of medium spiny neostriatal neurons (Petrálie and Wenthold, 1992) where they mediate a large proportion of the glutamate-induced excitation (Cepeda et al., 1993).

Considerable evidence indicates that AMPA receptors are regulated by protein phosphorylation. For example, activation of cAMP-dependent protein kinase (PKA) in hippocampal (Greengard et al., 1991; Wang et al., 1991; Rosenmund et al., 1994; Kameyama et al., 1998) or neostriatal (Yan et al., 1999) neurons increases AMPA currents recorded from these cells. The notion that these effects occur as the result of phosphorylation of the receptor is supported by the observation that the GluR1 subunit is phosphorylated in response to PKA activation in human embryonic kidney-293 cells (Blackstone et al., 1994; Tan et al., 1994) (but see McGlade-McCulloh et al., 1993). Moreover, a PKA-dependent residue, Ser⁸⁴⁵, is required for PKA-mediated potentiation of peak current carried by homomeric GluR1 channels (Roche et al., 1996). AMPA receptors also can be regulated by phosphorylation of Ser⁸³¹ after activation of calcium/calmodulin-

dependent kinase II (CaMKII) (Barria et al., 1997) or protein kinase C (PKC) leading to potentiation of AMPA currents in hippocampal neurons (Barria et al., 1997; Mammen et al., 1997). Whereas PKA-mediated phosphorylation at Ser⁸⁴⁵ enhances AMPA currents by increasing channel open time probability (Roche et al., 1996), phosphorylation at Ser⁸³¹ increases single-channel conductance of AMPA receptors (Derkach et al., 1999). Thus, PKA and CaMKII/PKC signaling cascades control distinct properties of the AMPA receptor by regulating unique GluR1 residues.

Dopamine and cAMP-regulated phosphoprotein *M_r* 32 kDa (DARPP-32); is a neostriatum-enriched substrate for PKA (Walaas et al., 1983; Ouimet et al., 1984). Phosphorylation at Thr³⁴ by PKA converts DARPP-32 into an efficient inhibitor of protein phosphatase 1 (PP1) (Hemmings et al., 1984), a serine/threonine phosphatase enriched in the dendritic spines of neostriatal neurons (Ouimet et al., 1995). Studies using mice lacking the gene for DARPP-32 indicate that DARPP-32 mediates many physiological and behavioral effects of dopamine (Fienberg et al., 1998). A recent study (Yan et al., 1999) showed that whole-cell AMPA currents are enhanced by D1 dopamine receptors via a pathway involving DARPP-32, PP1, and the PP1-targeting protein spinophilin (Allen et al., 1997). A principal goal of the present study was to determine whether regulation of neostriatal GluR1 subunit phosphorylation by this pathway is a plausible mechanism for the physiological control of AMPA receptors by dopamine.

Psychostimulants increase the synaptic availability of dopamine in the brain (Hyman, 1996; Koob and LeMoal, 1997), leading to both acute and long-term plastic changes in dopaminergic neurons (Nestler and Aghajanian, 1997). Some of these effects appear to require the activation of glutamate receptors, although the precise role that glutamate plays in addiction is unclear (Wolf, 1998). Thus, a second objective of this study was to investigate the

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Figure 1. Effects of protein kinase activators on GluR1 phosphorylation in neostriatal slices. Neostriatal slices were prepared from normal C57BL/6 mice and incubated for 5 min in the absence (*Control*) or presence of the adenylyl cyclase activator forskolin (*A*) or the PKC activator PDBu (*B*). Levels of phospho-Ser⁸⁴⁵ (*A*) or phospho-Ser⁸³¹ (*B*) were determined by SDS-PAGE and immunoblotting. Phospho-GluR1 levels were detected, quantitated by densitometry, and expressed as percentage of the level detected in control slices. Arrows (*top panels*) indicate the position of the phospho-GluR1 bands in representative autoradiograms. Data (*bottom panels*) are expressed as means $\pm$ SEM for three experiments (* p < 0.05 compared with control, Mann–Whitney *U* test).

possibility that regulation of AMPA receptor phosphorylation *in vivo* by psychostimulants might contribute to these effects.

MATERIALS AND METHODS

Preparation and treatment of neostriatal slices. Male C57BL/6 mice (8–12 weeks in age) were killed by decapitation. The brain was rapidly transferred to an ice-cold surface where it was blocked and fixed to the cutting surface of a Vibratome (Ted Pella) maintained at 4°C. The brain was placed in cold, oxygenated (95% O₂ and 5% CO₂) Krebs' bicarbonate buffer of the following composition (in mM): 125 NaCl, 5 KCl, 26 NaHCO₃, 1.5 CaCl₂, 1.5 MgSO₄, and 10 glucose, pH 7.4. Coronal slices of mouse brain (400 μ m in thickness) were cut and pooled in 10 ml of cold buffer. Neostriatal slices were cut from the coronal sections under a dissecting microscope. The slices were pooled, then transferred individually to 4 ml polypropylene tubes containing 2 ml of fresh, cold, oxygenated buffer. The tissue was preincubated for 15 min at 30°C, the buffer was replaced, and tissue sections were preincubated for an additional 30 min. At the end of this second preincubation period, the buffer was replaced with Krebs' buffer or buffer containing the indicated test substances for 30 sec to 60 min. After treatment, the slices were immediately frozen in liquid nitrogen and stored at –80°C until assayed.

In some experiments, neostriatal slices were prepared from C57BL/6 mice (8–10 weeks of age) lacking the gene for DARPP-32 (Fienberg et al., 1998). DARPP-32 knock-out mice and wild-type controls were generated from the offspring of heterozygous mating pairs. All mice were age-matched, and only males were used.

Preparation of GluR1 fusion protein for stoichiometric analysis of Ser⁸⁴⁵ and Ser⁸³¹ phosphorylation. A bacterial protein comprising the last putative intracellular domain of GluR1 (residues 809–889) fused to glutathione-S-transferase (GST) was prepared as follows: rat brain cDNA was used as a template for PCR amplification of the GluR1 sequence using the primers TGC GTC GAC CGA GTT CTG CTA CAA ATC CC; TTC GCG GCC GCT CCA GTT ACA ATC CTG TG (Operon). The resulting 272 bp fragment was digested with *SalI*–*NotI* and ligated in frame with the GST coding sequence contained in a

Figure 2. Effect of activation of D1-type or D2-type dopamine receptors on neostriatal GluR1 phosphorylation. Neostriatal slices were prepared from normal C57BL/6 mice. *A*, At time 0 slices were incubated in the absence or presence of the D1 receptor antagonist SCH23390 (1 μ M; D1 Antag) or the D2 receptor antagonist sulpiride (1 μ M; D2 Antag), as indicated. At 10 min slices were incubated in normal buffer alone (*Control*), with dopamine (DA) (100 μ M) plus the dopamine uptake inhibitor nomifensine (10 μ M), SKF81297 (1 μ M) (D1) or quinpirole (1 μ M) (D2), with or without SCH23390 or sulpiride for an additional 5 min. *B*, Slices were incubated with either normal buffer (*Control*) or with SKF81297 (1 μ M) or quinpirole (1 μ M) for 5 min. The solid arrows indicate the position of phospho-Ser⁸⁴⁵ GluR1 (*A*) or phospho-Ser⁸³¹ GluR1 (*B*). The open arrows indicate cross-reactive bands detected by the phosphorylation state-specific antibody. The intensity of these bands did not change as a function of dopamine agonist or antagonist treatment. Phospho-Ser⁸⁴⁵ GluR1 (*A*) or phospho-Ser⁸³¹ GluR1 (*B*) was detected in the samples, quantitated by densitometry, and expressed as percentage of the level in control slices. Data are presented as means $\pm$ SEM for three experiments (* p < 0.05 compared with control, Mann–Whitney *U* test).

bacterial expression plasmid pGex-4T-2 (Pharmacia, Piscataway, NJ). This plasmid was sequenced to ensure fidelity of PCR amplification and transformed into *Escherichia coli* strain BL21 DE3. Bacteria were grown at 37°C to an OD₆₀₀ of 0.5. The temperature was reduced to 30°C, and

expression was induced over 2 hr by the addition of isopropylthio- β -galactoside to a final concentration of 0.1 mM. Cells were harvested by centrifugation and lysed in a French press. After clarification of the lysate by centrifugation at $30,000 \times g$ for 20 min, the fusion protein was purified by affinity chromatography on glutathione-agarose beads (Pharmacia), using 5 mM glutathione and 50 mM Tris, pH 8.0, for elution.

GST-GluR1 (35 μ M) was preparatively phosphorylated under the following conditions: 10 μ g/ml purified PKA in 50 mM Tris, pH 8.0, 10 mM magnesium chloride, 0.4 mM EGTA, 50 μ M [γ - 32 P] ATP or 1 μ g/ml purified CaMKII in 50 mM Tris, pH 8.0, 10 mM magnesium chloride, 0.4 mM EGTA, 2 mM DTT, 1 μ M calmodulin, 1.5 mM calcium chloride, and 50 μ M [γ - 32 P]ATP. Reactions were stopped by the addition of Laemmli buffer. Stoichiometry of the phosphorylated fusion proteins was calculated based on PhosphorImager (Molecular Dynamics, Eugene, OR) analysis of phosphate incorporation. Known quantities of these proteins were compared with tissue samples, the levels of phospho-GluR1 and GluR1 were detected by ECL, and the resulting values used to calculate stoichiometries of phosphorylation for the Ser⁸⁴⁵ and Ser⁸³¹ sites in neostriatal tissue.

Drug treatment and microwave irradiation of mice. Male C57BL/6 mice (8–12 weeks in age) were injected with vehicle (0.9% NaCl in water) or vehicle containing various concentrations of either methamphetamine HCl (20 or 30 mg/kg, s.c.) or cocaine HCl (10 or 20 mg/kg, i.p.). To insure preservation of phosphoproteins in their *in vivo* phosphorylation state, the animals were killed at various times (15–60 min) after injection by focused microwave irradiation (4.5–5 kW for 1.4 sec) using a small-animal microwave (Murimachi Kikai, Tokyo, Japan). The brains were rapidly removed, and the neostriatum was dissected and stored at -80°C until assayed for phosphoprotein levels.

Immunoblotting for phospho-GluR1. Frozen tissue samples were sonicated in 1% SDS. Small aliquots of the homogenate were retained for protein determination by the BCA protein assay method (Pierce, Rockford, IL) using bovine serum albumin as a standard. Equal amounts of protein (50 μ g for slice experiments; 250 μ g for microwave experiments) were loaded onto 7.5% acrylamide gels. The proteins were separated by SDS-PAGE (Laemmli, 1970), and transferred to nitrocellulose membranes (0.2 μ m) (Schleicher and Schuell) by the method of Towbin et al. (1979). Membranes were blocked for 30–60 min in PBS (in mM: 124 NaCl, 4 KCl, 10 Na_2HPO_4 , and 10 KH_2PO_4 , pH 7.2) containing 5% nonfat dry milk and 0.2% Tween 20 (Blotto). The membranes were immunoblotted using antisera that selectively detect either the Ser⁸⁴⁵-phosphorylated or the Ser⁸³¹-phosphorylated form of GluR1 (1:200 dilutions for each antibody) (Kameyama et al., 1998), or an antiserum (PharMingen, San Diego, CA; 1:10,000 dilution) that detects the C-terminal region of GluR1, irrespective of phosphorylation state. In some experiments these samples were also immunoblotted with monoclonal antibody 23 (1:750 dilution) (Snyder et al., 1992), a phosphorylation state-specific monoclonal antibody raised against a DARPP-32 peptide containing phospho-Thr³⁴, the site phosphorylated by PKA.

Antibody binding was revealed by incubation with either a goat anti-rabbit horseradish peroxidase-linked IgG or a goat anti-mouse horseradish peroxidase-linked IgG (each at a 1:6000–8000 dilution) (Pierce) and the ECL immunoblotting detection system (Amersham, Arlington Heights, IL). Chemiluminescence was detected by autoradiography using DuPont NEN (Boston, MA) autoradiography film, and bands were quantified by analysis of scanned images by NIH Image 1.52 software. Because the linear range for quantitation of ECL signals by densitometry is limited, several film exposures were obtained for each set of samples to insure that signals were within a density range that allowed accurate quantitation.

In all of the experiments for this study, nitrocellulose membranes were sequentially analyzed for phospho-Ser⁸⁴⁵ GluR1 or for phospho-Ser⁸³¹ GluR1, and then for total levels of C-terminal GluR1. After probing a membrane for phospho-GluR1, the filter was washed three times for 5 min each in PBS to remove any remaining chemiluminescent reagent. The membrane was then stripped of antibody by incubation at 60°C for 60 min in a buffer containing 100 mM 2-mercaptoethanol, 2% SDS, and 62.5 mM Tris/HCl, pH 6.7. The filter was then washed several times in large volumes of PBS and blocked in Blotto for 30–60 min before immunoblotting with the C-terminal GluR1 antibody.

Data were analyzed by Student's *t*-test or Mann–Whitney *U* test, as indicated, with significance defined as $p < 0.05$.

A. Ser⁸⁴⁵

B. Ser⁸³¹

Figure 3. Effects of protein phosphatase inhibitors on the phosphorylation state of neostriatal GluR1 at Ser⁸⁴⁵ and Ser⁸³¹. Mouse neostriatal slices were incubated for 60 min in the absence (Control) or presence of either okadaic acid (1 μ M) (OKA) or calyculin A (0.5 μ M) (Cal A), inhibitors of PP1/PP2A, or cyclosporin A (5 μ M) (Cyc A), an inhibitor of PP2B. Ser⁸⁴⁵ phospho-GluR1 (A) and Ser⁸³¹ phospho-GluR1 (B) were detected, quantitated by densitometry, and expressed as percentage of the level in control slices. Data are expressed as means $\pm$ SEM for three experiments (* $p < 0.05$ compared with control, Mann–Whitney *U* test).

RESULTS

Differential phosphorylation of GluR1 in neostriatum by protein kinase activators

GluR1 is phosphorylated *in vitro* at Ser⁸⁴⁵ by PKA (Blackstone et al., 1994) and at Ser⁸³¹ by PKC (Mammen et al., 1997) or CaMKII (McGlade-McCulloh et al., 1993). Phosphorylation of GluR1 at Ser⁸⁴⁵, in response to activation of PKA, potentiates AMPA currents by increasing open time probability of the receptor channel (Roche et al., 1996). The phosphorylation of GluR1 at Ser⁸³¹ in response to activation of CaMKII or PKC increases AMPA receptor function by enhancing channel conductance (Derkach et al., 1999). Treatment of neostriatal slices with forskolin (50 μ M), an adenylyl cyclase activator, increased phosphorylation of Ser⁸⁴⁵ approximately eightfold ($776 \pm 20\%$ of control in three experiments) (Fig. 1A). The PKC activator phorbol 12,13-dibutyrate (PDBu) (5 μ M) (Fig. 1B) increased the phosphorylation of GluR1 at Ser⁸³¹ by an average of 10-fold ($1005 \pm 28\%$ of control in three experiments) in neostriatal slices.

Figure 4. Role of DARPP-32 in the dopamine-induced phosphorylation of neostriatal GluR1 on Ser⁸⁴⁵. *A*, Mouse neostriatal slices prepared from wild-type mice (left) or DARPP-32 knock-out mice (right) were incubated in the absence or presence of dopamine (100 μ M) plus nomifensine (10 μ M) for the indicated times. Samples were immunoblotted for Ser⁸⁴⁵-phosphorylated GluR1. The level of phospho-GluR1 was quantitated by densitometry, and the data were expressed as percentage of either the wild-type or knock-out control level. The results are expressed as means $\pm$ SEM for five experiments (* p < 0.05 compared with 0 time, Mann–Whitney U test). *B*, Slices from wild-type or knock-out mice were incubated with a maximally effective concentration of the PP1/2A inhibitor, okadaic acid (OKA; 1 μ M) for 60 min. The level of phospho-Ser⁸⁴⁵ GluR1 was quantitated by densitometry, and the data were expressed as percentage of the wild-type control level. Results are expressed as means $\pm$ SEM for three experiments (* p < 0.05 compared with control, Mann–Whitney U test).

Phosphorylation state-specific antibodies were used to compare the levels of phospho-Ser⁸⁴⁵ and phospho-Ser⁸³¹ GluR1 in neostriatal slices with known quantities of GluR1 fusion proteins phosphorylated by PKA or CaMKII, respectively. A low, basal stoichiometry of phosphorylation at Ser⁸⁴⁵ was measured (~0.05 mol/mol) in slices, which was increased to ~0.33 mol/mol after 5 min of forskolin treatment (50 μ M). Ser⁸³¹ was also phosphory-

lated at a low level in untreated slices (a stoichiometry of 0.06 mol/mol). The stoichiometry of phosphorylation at Ser⁸³¹ was increased to ~0.26 mol/mol after 5 min of PDBu treatment (5 μ M). These data indicate that GluR1 is phosphorylated in intact neostriatal cells at Ser⁸⁴⁵ or Ser⁸³¹, after activation of either PKA or PKC/CaMKII, respectively.

Phosphorylation of GluR1 in response to dopamine receptor stimulation

Dopamine receptors are subdivided into a D1 class (i.e., D1, D5 subtype receptors) and a D2 class (i.e., D2, D3, and D4 subtype receptors) with the D1 and D2 subtypes being the most highly enriched in mammalian neostriatum (Sibley and Monsma, 1992). D1 class receptors stimulate adenylyl cyclase and increase cAMP formation (Stoof and Kebabian, 1981). D2 class receptors, via interactions with specific G-proteins, can be coupled to multiple effector systems, including adenylyl cyclase, Ca²⁺ and K⁺ channels, and phospholipase C (Huff, 1996). To evaluate the potential contribution of D1-subtype and D2-subtype dopamine receptors to the regulation of GluR1 phosphorylation, we measured the effects of dopamine and selective D1 and D2 agonists and antagonists on phosphorylation at Ser⁸⁴⁵ and Ser⁸³¹.

Dopamine increased the phosphorylation of GluR1 at Ser⁸⁴⁵ (428 $\pm$ 68% compared with control) (Fig. 2*A*). Levels of phosphorylation were maximal at between 2.5 and 5 min, and declined to basal values by 10 min of incubation with dopamine (data not shown). The effect of dopamine was blocked by pretreatment of slices with a D1-type receptor antagonist, SCH23390 (1 μ M) (127 $\pm$ 21% compared with control). In contrast, the D2-type receptor antagonist sulpiride (1 μ M) had no effect on dopamine-stimulated phosphorylation of GluR1 (565 $\pm$ 81% compared with control) or on D1 agonist-induced phosphorylation of GluR1 (data not shown). Neither the D1 nor the D2 antagonist alone had a significant effect on the basal level of GluR1 phosphorylation (data not shown) indicating that any endogenously released dopamine was not exerting a significant regulation of this site. The phosphorylation of Ser⁸⁴⁵ was increased by treatment of slices with SKF81297, a D1 agonist (337 $\pm$ 82% of control; Fig. 2*A*), whereas quinpirole, a D2 agonist, had no effect on basal levels (91 $\pm$ 12% of control; Fig. 2*A*). Quinpirole did not reduce phosphorylation of Ser⁸⁴⁵ induced by a D1 agonist (data not shown). These data indicate that dopamine-induced phosphorylation of GluR1 is most likely mediated by activation of a D1-receptor/PKA-dependent signaling pathway.

D2 agonists have recently been shown to increase intracellular Ca²⁺ levels in dissociated neostriatal cells (Hernandez et al., 1999), an effect that could result in activation of Ca²⁺-dependent kinases such as CaMKII. Thus, we evaluated the effects of D1 and D2 receptor activation on Ser⁸³¹ phosphorylation. In contrast to the robust increase in phosphorylation of Ser⁸⁴⁵ observed in response to dopamine or a D1-selective agonist (Fig. 2*A*), neither dopamine, SKF81297 (106 $\pm$ 11% of control), quinpirole (82 $\pm$ 12% of control), nor SKF81297 plus quinpirole (107 $\pm$ 15% of control) had a significant effect on Ser⁸³¹ phosphorylation (Fig. 2*B*).

Regulation of Ser⁸⁴⁵ phosphorylation: involvement of a DARPP-32/PP1 cascade

The ability of serine/threonine protein phosphatases to regulate the basal phosphorylation of GluR1 in neostriatal cells was examined. Neostriatal slices were incubated with either calyculin A (500 nM) or okadaic acid (1 μ M), inhibitors of both PP1 and

Figure 5. Effect of acute cocaine on neostriatal GluR1 phosphorylation at Ser⁸⁴⁵ and Ser⁸³¹ *in vivo*. Normal C57Bl/6 mice were injected (intraperitoneally) with vehicle alone (*Saline*) or with vehicle containing cocaine (20 mg/kg) and killed by focused microwave irradiation at 15, 30, or 60 min after injection. Neostriatum was dissected from each brain and analyzed for phospho-Ser⁸⁴⁵ and phospho-Ser⁸³¹ GluR1. The arrows indicate the position of GluR1, as verified by immunoblotting with an antibody against a C-terminal sequence of GluR1. **A**, Representative experiment showing immunoblots of phospho-Ser⁸⁴⁵ GluR1 from three individual animals for each treatment condition. **B**, Quantitation of phospho-Ser⁸⁴⁵ and phospho-Ser⁸³¹ in three experiments, each analyzed in triplicate (* $p < 0.05$, compared with saline; Mann–Whitney *U* test).

protein phosphatase 2A (PP2A). Calyculin A increased phospho-Ser⁸⁴⁵ GluR1 to a level $600 \pm 129\%$ of control after 60 min of incubation (Fig. 3A). Similarly, okadaic acid increased phospho-Ser⁸⁴⁵ GluR1 to a maximal level of $783 \pm 102\%$ of control after 60 min. By comparison, cyclosporin A, an inhibitor of calcineurin (protein phosphatase 2B, PP2B), had no significant effect on the basal level of phosphorylated Ser⁸⁴⁵ GluR1 after a 60 min incubation period ($104 \pm 4\%$ compared with control; Fig. 3A) or on the state of phosphorylation of DARPP-32 at Thr³⁴ (data not shown). Phosphorylation of GluR1 at Ser⁸³¹ was slightly increased by calyculin A ($256 \pm 78\%$ of control) in contrast to the absence of effect of okadaic acid ($137 \pm 36\%$ of control) or cyclosporin A ($104 \pm 25\%$ of control) (Fig. 3B). However, the effect of calyculin A was not statistically significant ($p > 0.05$; Mann–Whitney *U* test, for three experiments). These data indicate that Ser⁸⁴⁵ is a better substrate than Ser⁸³¹ for regulation by PP1 and/or PP2A in neostriatal neurons.

To investigate the specific contribution of PP1 to dopamine-induced phosphorylation of GluR1, we compared Ser⁸⁴⁵ phosphorylation in wild-type mice and in mice bearing a targeted deletion of the gene for the selective PP1 inhibitor DARPP-32 (Fienberg et al., 1998). These experiments revealed that there was no significant difference in the basal phosphorylation of Ser⁸⁴⁵ as the result of the DARPP-32 knock-out. Dopamine (100 μ M) induced a threefold to fourfold increase in phosphorylation of GluR1 at Ser⁸⁴⁵ in neostriatal slices from wild-type mice ($324 \pm 85\%$ of control at 2.5 min and $383 \pm 26\%$ of control at 5 min), but not from DARPP-32 knock-out mice ($94 \pm 41\%$ of control at 2.5 min and $174 \pm 72\%$ of control at 5 min; Fig. 4A). A similar attenuation of dopamine-induced phosphorylation was observed in nucleus accumbens slices (data not shown). These data indicate that inhibition of PP1 is required for dopamine-induced increases in GluR1 phosphorylation. In support of this conclusion, we found that direct, pharmacological inhibition of PP1/PP2A affected GluR1 phosphorylation comparably in wild-type and knock-out slices. In this series of experiments okadaic acid increased Ser⁸⁴⁵ phosphorylation in slices from both wild-type and DARPP-32 knock-out mice ($878 \pm 210\%$ of control in wild-type mice and $1008 \pm 50\%$ of control in knock-out mice) after 60 min of incubation (Fig. 4B).

Regulation of GluR1 phosphorylation by cocaine *in vivo*

To evaluate the possible involvement of psychostimulants in the control of AMPA receptor phosphorylation *in vivo*, intact mice were treated with drugs that are known to enhance dopamine availability in the neostriatum, and the effect on the phosphorylation of GluR1 was examined. We first measured the effect of cocaine (20 mg/kg, i.p.), a drug that increases dopamine availability through blockade of the dopamine transporter, on the phosphorylation of GluR1 at Ser⁸⁴⁵. Cocaine administration produced a behavioral activation in mice characterized by increased locomotor behavior that was evident within 15 min of injection. Cocaine also caused a large increase in the phosphorylation state of GluR1 at Ser⁸⁴⁵ in neostriatum, as observed 15 min (saline, $100 \pm 18\%$; cocaine, $612 \pm 198\%$) and 30 min (saline, $100 \pm 22\%$; cocaine, $604 \pm 170\%$) after injection (Fig. 5). The effect of cocaine treatment on GluR1 phosphorylation rapidly dissipated, as Ser⁸⁴⁵ phosphorylation returned to the basal level within 60 min (saline, $100 \pm 4\%$; cocaine, $89 \pm 25\%$) of drug administration (Fig. 5). In contrast, cocaine treatment did not significantly affect the level of phosphorylation of GluR1 at Ser⁸³¹ in neostriatum compared with time-matched saline-injected controls at any time point examined ($89 \pm 30\%$ of control at 15 min; $80 \pm 18\%$ of control at 30 min; and $98 \pm 8\%$ of control at 60 min) (Fig. 5). These data indicate that acute cocaine injection selectively regulates phosphorylation of GluR1 at Ser⁸⁴⁵.

Role of DARPP-32 in cocaine-mediated regulation of Ser⁸⁴⁵ phosphorylation *in vivo*

It seemed possible that cocaine-stimulated dopamine release could increase GluR1 phosphorylation by at least two mechanisms. First, dopamine and D1-receptor-mediated activation of PKA would be expected to lead directly to phosphorylation of Ser⁸⁴⁵. Second, PKA-mediated phosphorylation of DARPP-32 at Thr³⁴ would be expected to inhibit PP1 activity, leading to an increase in phosphorylation state of PP1 substrates, including GluR1. Consistent with this idea, cocaine did induce a severalfold increase in phospho-Thr³⁴ DARPP-32 ($605 \pm 205\%$ of control) (Fig. 6A), concurrent with the increase in Ser⁸⁴⁵ phosphorylation in wild-type mice. A possible role for DARPP-32 in the *in vivo*

A. 10 mg/kg cocaine

B. 20 mg/kg cocaine

Figure 6. Effect of acute cocaine on *in vivo* phosphorylation of GluR1 at Ser⁸⁴⁵: comparison of wild-type and DARPP-32 knock-out mice. Wild-type or DARPP-32 knock-out mice were injected (intraperitoneally) with saline vehicle (Sal) or with (10 mg/kg) (A) or (20 mg/kg) (B) cocaine (Coc) and killed by focused microwave irradiation 30 min later. The levels of phospho-Ser⁸⁴⁵ GluR1 and phospho-Thr³⁴ DARPP-32 were detected in neostriatum by immunoblotting, and the data were quantitated by densitometry. Arrows indicate the position of phospho-GluR1 and phospho-DARPP-32 bands. **A, Top.** An autoradiogram of a representative experiment shows the level of phospho-GluR1 in saline-injected or cocaine-injected wild-type and DARPP-32 knock-out mice. The bar graph summarizes phospho-Ser⁸⁴⁵ GluR1 levels in the neostriatum of the wild-type and knock-out mice. Data are expressed as means $\pm$ SEM for

phosphorylation of GluR1 was further evaluated by measuring cocaine-induced GluR1 phosphorylation in DARPP-32 knock-out mice. The level of GluR1 expressed in neostriatum was unaffected by the genetic deletion of DARPP-32 (data not shown). However, the increase in GluR1 phosphorylation at Ser⁸⁴⁵ in neostriatum observed after treatment with a low dose of cocaine (10 mg/kg, i.p.) ($671 \pm 75\%$ of control in this series of experiments) was greatly attenuated in mice lacking DARPP-32 ($138 \pm 18\%$ of control; Fig. 6A).

A previous report from this laboratory (Fienberg et al., 1998) showed that the effect of DARPP-32 knock-out on cocaine-induced locomotor behavior was dose-dependent. Thus, the increase in locomotor behavior elicited by a low (10 mg/kg), but not by a high (20 mg/kg) dose of cocaine was blocked in DARPP-32 knock-out mice. We examined the dose dependency of the effects of cocaine on GluR1 phosphorylation. A 20 mg/kg concentration of cocaine induced an approximately sevenfold increase in Ser⁸⁴⁵ phosphorylation ($709 \pm 198\%$ of control; Fig. 6B) in wild-type mice. This increase was not significantly reduced in DARPP-32 knock-out mice ($479 \pm 165\%$ of control; Fig. 6B).

Regulation of GluR1 phosphorylation by methamphetamine *in vivo*

Methamphetamine-HCl, a substituted analog of D-amphetamine with profound abuse potential, also increases dopamine availability in the neostriatum. The effect of systemic injection of methamphetamine on the *in vivo* phosphorylation of GluR1 in mouse neostriatum was examined. Injection of mice with methamphetamine (i.e., 20 or 30 mg/kg, s.c.) induced behavioral activation, eliciting intense grooming behavior and hypersensitivity to touch. Treatment of mice with methamphetamine (20 mg/kg, s.c.) increased phospho-Ser⁸⁴⁵ levels by twofold to threefold in neostriatum 30 min after drug injection ($254 \pm 32\%$ of saline-injected wild-type) (Fig. 7B). In contrast, methamphetamine treatment had no effect on the level of phospho-Ser⁸³¹ ($93 \pm 12\%$ of saline-injected control) (Fig. 7D). This dose of methamphetamine also increased the phosphorylation of DARPP-32 on Thr³⁴ by more than twofold in the neostriatum of mice ($270 \pm 82\%$ of control) (Fig. 7E). The increase in phospho-Ser⁸⁴⁵ induced by this concentration of methamphetamine was fully blocked in DARPP-32 knock-out mice ($120 \pm 28\%$ of saline-injected wild type) (Fig. 7B). A higher concentration of methamphetamine (30 mg/kg, s.c.) induced a fourfold to fivefold increase in Ser⁸⁴⁵ phosphorylation in the neostriatum of wild-type mice compared to vehicle-treated controls ($486 \pm 80\%$ of saline-injected wild types) (Fig. 7C). The higher concentration of methamphetamine induced a significant increase in Ser⁸⁴⁵ phosphorylation even in the DARPP-32 knock-out mice ($325 \pm 40\%$ of saline-injected knock-out mice) (Fig. 7C). The level of Ser⁸⁴⁵ phosphorylation that was observed in the knock-out mice, however, was significantly reduced compared to wild-type mice (Fig. 7C).

eight mice per group (* $p < 0.01$ compared with saline-injected mice; Student's *t* test). **A, Bottom.** A typical autoradiogram shows the level of phospho-DARPP-32 in a saline-injected and a cocaine-injected wild-type mouse neostriatum. The bar graph shows phospho-Thr³⁴ DARPP-32 levels in the neostriata of mice. Data are expressed as means $\pm$ SEM for four mice per group (* $p < 0.05$ compared with saline-injected mice; Mann-Whitney *U* test). **B.** Data are expressed as means $\pm$ SEM for 3–5 mice per group (* $p < 0.05$ compared with respective saline-injected controls; Mann-Whitney *U* test).

Figure 7. Effect of acute methamphetamine administration on *in vivo* phosphorylation of GluR1 at Ser⁸⁴⁵ and Ser⁸³¹ in wild-type and DARPP-32 knock-out mice. Wild-type (*A–E*) or DARPP-32 knock-out (*B, C*) mice were injected with vehicle or methamphetamine (*Meth*) 20 mg/kg (subcutaneously) (*B, D, E*) or 30 mg/kg (*A, C*) and killed by focused microwave irradiation 30 min later. Neostriatal phospho-Ser⁸⁴⁵ GluR1 (*A–C*) and phospho-Ser⁸³¹ GluR1 (*A, D*) and phospho-Thr³⁴ DARPP-32 (*E*) were detected by immunoblotting and quantitated by densitometry. Data are expressed as percentage of values for saline-injected wild-type mice and represent means $\pm$ SEM for nine (*B, C*), three (*D*), or eight animals (*E*) per group (* p < 0.05 compared with saline-injected mice; † p < 0.05, compared with wild-type *Meth* alone; Student's *t* test).

DISCUSSION

Using neostriatal slices, we have demonstrated that dopamine stimulates phosphorylation of GluR1 at Ser⁸⁴⁵ through activation of D1-type dopamine receptors and stimulation of adenylyl cyclase and PKA. We have also demonstrated that psychostimulants rapidly and reversibly increase the phosphorylation of GluR1 at Ser⁸⁴⁵ in the neostriatum *in vivo*. Estimates of the stoichiometry of GluR1 phosphorylation indicate that a substantial proportion of the GluR1 is phosphorylated at Ser⁸⁴⁵ in response either to dopamine in slices or to cocaine and methamphetamine *in vivo*. These data indicate that regulation of the state of phosphorylation of GluR1 represents a probable mechanism for control of AMPA receptor currents by dopamine and psychostimulants. Results from a previous study involving a Ser⁸⁴⁵-Ala mutation of GluR1 indicated that phosphorylation of GluR1 at Ser⁸⁴⁵ was required to potentiate peak currents carried by homomeric GluR1 receptors (Roche et al., 1996). In addition, PKA agonists reversed synaptic depression of AMPA currents in hippocampal slices by increasing phosphorylation at Ser⁸⁴⁵ (Kameyama et al., 1998; Lee et al., 1998). Taken together, these data suggest that dopamine-induced, PKA-mediated, phosphorylation of Ser⁸⁴⁵ in slices and *in vivo* is likely to enhance glutamatergic transmission through AMPA receptors.

AMPA receptor currents are also modulated by phosphorylation of GluR1 at Ser⁸³¹ (Barria et al., 1997; Derkach et al., 1999). This residue, which is phosphorylated *in vitro* by PKC or CaMKII (McGlade-McCulloh et al., 1993; Blackstone et al., 1994), is also phosphorylated to a low but measurable stoichiometry in untreated neostriatum. However, the phosphorylation of this site was not regulated in response to dopaminergic activity in slices or *in vivo* with either wild-type mice (Fig. 2*B*) or in DARPP-32 knock-out mice (data not shown). The data further suggest that dopamine and D1 agonists regulate AMPA receptor currents through the selective phosphorylation of Ser⁸⁴⁵. The proportion of GluR1 subunits phosphorylated at Ser⁸³¹ is greatly increased by activation of PKC (Fig. 1*B*) or CaMKII signaling pathways

(our unpublished observations), suggesting that both Ser⁸⁴⁵ and Ser⁸³¹ can serve to mediate neurotransmitter-specific effects on AMPA receptors.

Dopamine-induced phosphorylation at Ser⁸⁴⁵ in slices and psychostimulant-induced phosphorylation of Ser⁸⁴⁵ *in vivo* were reduced in the neostriatum of DARPP-32 knock-out mice. These data are consistent with the loss of D1-mediated enhancement of whole-cell AMPA currents in neostriatal neurons from DARPP-32 knock-out mice (Yan et al., 1999). Taken together, the results indicate that a pathway involving PKA and DARPP-32/PP1 functionally regulates AMPA receptors in basal ganglia neurons. It is unlikely that the deficits in psychostimulant action seen in the absence of DARPP-32 are attributable to a reduced ability to release dopamine. This conclusion is supported by a recent study in which dopamine overflow in neostriatum was measured by amperometry in response to stimulation of the medial forebrain bundle. Stimulation-evoked dopamine release *in vivo* was found to be equivalent in wild-type and DARPP-32 knock-out mice (F. Gonon, A. Fienberg, and P. Greengard, unpublished observations). In addition, deletion of the DARPP-32 gene does not significantly affect either the density of D1 receptors or their affinity for dopamine ligands (P. Svenningsson, B. Fredholm, A. Fienberg, and P. Greengard, unpublished observations). Moreover, Ser⁸⁴⁵ phosphorylation is increased comparably in slices from wild-type and DARPP-32 knock-out mice in response either to forskolin, which induces PKA activity (G. Snyder, unpublished observations), or to drugs that inhibit PP1/PP2A activity (Fig. 4*B*). Together, the results indicate that the loss of GluR1 regulation is caused by a loss of ability to regulate PP1 activity.

The phosphorylation of Ser⁸⁴⁵ elicited by either cocaine or methamphetamine is blocked in DARPP-32 knock-out mice at low but not at high concentrations of these drugs. These data are consistent with previous reports (Fienberg et al., 1998; Snyder et al., 1998) showing that DARPP-32 knock-out blocks responses to low (physiological) levels of activation of transmission, but not to

high (often supraphysiological) levels of activation. For example, cocaine-induced locomotor activation in mice induced by a 10 mg/kg dose of drug is fully blocked in DARPP-32 knock-out mice, whereas activation induced by a 20 mg/kg cocaine injection is unaffected by DARPP-32 knock-out (Fienberg et al., 1998).

Taken together, we propose the following interpretation of these data: that PKA increases phosphorylation of Ser⁸⁴⁵ by at least two mechanisms: (1) by directly phosphorylating Ser⁸⁴⁵, and (2) by increasing phosphorylation of DARPP-32 at Thr³⁴, leading to inhibition of PP1 activity. Thus, our data suggest that high concentrations of psychostimulants (i.e., 30 mg/kg methamphetamine or 20 mg/kg cocaine) provide sufficient activation of PKA to sustain PKA-mediated phosphorylation of GluR1 in the absence of PP1 inhibition. The results obtained at the lower concentrations of psychostimulants support a role for PKA-dependent phosphorylation of DARPP-32 at Thr³⁴ as an obligatory component of pathways that mediate a variety of biochemical, physiological, and behavioral effects of dopamine in neostriatal neurons (Fienberg et al., 1998; Greengard et al., 1999). However, a recent study (Bibb et al., 1999) has demonstrated that DARPP-32 is phosphorylated at Thr⁷⁵ by a cyclin-dependent kinase family member, cdk5, converting the protein into a PKA inhibitor. It will be important in future studies to evaluate the possible contribution of the regulation of Thr⁷⁵ of DARPP-32 to the control of AMPA channels by dopamine and psychostimulants.

Psychostimulants are known to regulate various biochemical indices of glutamate function including glutamate release, metabolism, and receptor expression (Wolf, 1998). Most of these changes do not occur in response to acute drug use, but appear only after repeated drug presentation, indicating that they represent indirect, long-term adaptive responses. The present results show that the regulation of AMPA receptor current through phosphorylation is likely to be a rapid, acute, and dramatic response to neostriatal dopamine release by psychostimulant drugs. The enhancement of AMPA receptor currents by these drugs represents an attractive mechanism by which glutamate receptors and their signaling pathways might be recruited to mediate a variety of effects, such as increased immediate early gene (IEG) expression (Konradi et al., 1996), which may eventually lead to sensitization. In support of this idea, AMPA or NMDA receptor antagonists block amphetamine-induced IEG expression (Konradi et al., 1996) and prevent the development of behavioral sensitization to amphetamine (Karler et al., 1989; Wolf, 1998).

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